IN THE CLAIMS

1. (Original) Use of compounds of the general Formula

3 (wherein

4 R3 stands for hydrogen or hydroxy;

s R1 stands for hydrogen or alkyl; and

6 R2 stands for alkyl)

and pharmaceutically acceptable acid addition salts for the

preparation of pharmaceutical compositions having neuroprotective

effect.

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2. (Original) Use according to Claim 1 for the

preparation of pharmaceutical compositions suitable for the

reduction of the consequences of acute ischemic or traumatic brain

and spinal damages, especially the various types of stroke or

cerebral vasospasm, severe brain vessel occlusion, neuronal loss

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- and its functional consequences in the case of head and spinal 6 injuries caused by accidents.
- 3. (Original) Use according to Claim 1 for the 1 preparation of pharmaceutical compositions having chronical 2 neurodegenerative effect. 3
- 4. (Original) Use according to Claim 3 for the 1 preparation of pharmaceutical composition suitable for the 2 treatment of motoneuron disease (ALS), sclerosis multiplex or 3 Creutzfeld- Jakob disease. 4
- 5. (Original) Use according to any of Claims 1-4 Claim 1 1 trirnethyl-bicyclo[2.2.1]heptane (deramciclane) or a pharmaceutically acceptable acid addition salt is used as compound of the general Formula I.
- 6. (Original) Use according to Claim 5 wherein 1 (1R,2S,4R)-(-)-2-(2-dimethylaminoethoxy)-2-phenyl-1,7,7-trimethyl-2 bicyclo[2.2.1]heptane-fumarate (deramciclane-fumarate) is used as 3 compound of the general Formula I.

- 7. (Original) Use according to Claim 1 wherein
- (1R, 2S, 4R) (-) 2 (2 dimethylaminoethoxy) 2 phenyl 1, 7, 7 trimethylbi
- cyclo[2.2.1]heptane of the Formula

- or a pharmaceutically acceptable acid addition salt containing not
- more than 0.2 % of (1R,3S,4R)-342-(N,Ndimethylaminoethyl)]-
- 7 1,7,7-timethyl-bicyclo[2.2.1]heptane-2- one of the Formula

- or a pharmaceutically acceptable acid addition salt thereof is used
- as compound of the general Formula I.

- 8. (Original) Use according to Claim 7 wherein
 (1R,28,4R)-(+2-(2-dimethylaminoethoxy)-2-phenyl-1,7,7-trimethylbicy
 clo[2.2.1]heptane-fumarate containing not more than 0.2 % of
 (1R,3S,4R)-342-(N,N-dimethylaminoethyl)]-1,7,7-trimethylbicyclo[2.2.1]heptane-2-one-fumarate is used as compound of the general
 Formula I.
- 9. (Currently amended) Use according to claim 1 any of
 claims 1-4 wherein
 (1R,2S,4R)-(-)-2-(2-methylaminoethoxy)-2-pheny1-1,7,7trimethyl-bicyclo[2.2.1]heptane;
 (1R,2S,7R)-2-pheny1-2-(2-methylamimethoxy)-7hydroxymethyl-1,7-dirnethyl-bicyclo[2.2.1]heptane; or
 (1R,2S,7R)-2-pheny1-2-(2-ethylaminoethoxy)-7hydroxymethyl-1,7-dimethyl-bicyclo[2.2.1]heptane
 or a pharmaceutically acceptable acid addition salt thereof is used
- or a pharmaceutically acceptable acid addition salt thereof is used as compound of the general Formula I.
- 10. (Currently amended) Neuroprotective pharmaceutical
 2 composition comprising as active ingredient a compound of the
 3 general Formula I as defined in claim 1 (wherein R', R2 and R3 are
 4 as stated in Claim 1) or a pharmaceutically acceptable acid
 5 addition salt thereof in admixture with inert pharmaceutically
 6 acceptable solid or liquid pharmaceutical active ingredient and/or
 7 auxiliary agent.

- 11. (Original) Pharmaceutical composition according to
 2 Claim 10 suitable for the reduction of the consequences of acute
 3 ischemic or traumatic brain and spinal damage, especially the
 4 various types of stroke or cerebral vasospasm, severe brain vessel
 5 occlusion, neuronal loss and its functional consequences in the
 6 case of head and spinal injuries caused by accidents.
- 12. (Original) Pharmaceutical composition according to
 2 Claim 10 suitable for the treatment of neurodegenerative diseases.
- 13. (Original) Pharmaceutical composition according to
 2 Claim 11 suitable for the treatment of motoneuron disease (ALS),
 3 sclerosis multiplex or Creutzfeld-Jakob disease.
- 14. (Currently amended) Pharmaceutical composition

 according to any of Claim 10-13 claim 1 comprising

 (1R,2S,4R)-(+27(2-dimethylaminoethoxy)-2-phenyl-1,7,7-trimethyl-bic

 yclo[2.2.1]heptane of the Formula II or a pharmaceutically

 acceptable acid addition salt as compound of the general Formula I.
- 15. (Original) Pharmaceutical composition according to
 2 Claim 14 comprising (1R,2S,4R)-(-)-2-(2-dimethylaminoethoxy)-23 phenyl-1,7,7-trimethyl-bicyclo[2.2.1]heptane-fumarate as compound
 4 of the general Formula I.

- 1 16. (Currently amended) Use of compounds of the general
 2 Formula I as defined in claim 1 (wherein R', R2 and R3 are as
 3 stated in Claim 1) and pharmaceutically acceptable acid addition
 4 salts thereof as neuroprotective pharmaceutical active ingredient.
- 17. (Original) Use according to Claim 16 for the
 reduction of the consequences of acute ischemic or traumatic brain
 and spinal damages, especially the various types of stroke or
 cerebral vasospasm, severe brain vessel occlusion, neuronal loss
 and its functional consequences in the case of head and spinal
 injuries caused by accidents.
- 18. (Original) Use according to Claim 16 for the treatment of chronical neurodegenerative diseases.
- 19. (Original) Use according to Claim 16 for the
 treatment of motoneuron disease (ALS), sclerosis multiplex or
 CreutzfeldJakob disease.

- 20. (Currently amended) Use of 1 (1R, 2S, 4R) - (-) - 2 - (2 - dimethylaminethoxy) - 2 - phenyl - 1, 7, 7 - trimethyl - bi2 cyclo[2.2.1]heptane of the Formula II as defined in claim 7 and 3 pharmaceutically acceptable acid addition salts thereof in the treatment of neuroprotective disorders, for the reduction of the 5 consequences of acute ischemic or traumatic brain and spinal damages, especially the various types of stroke or cerebral vasospasm, severe brain vessel occlusion, neuronal loss and its functional consequences in the case of head and spinal injuries caused by accidents, for the treatment of chronic neurodegenerative 10 diseases or for the treatment of motoneuron disease (ALS), 11 sclerosis multiplex or CreutzfeldJakob disease indications 12 according to Claims 16-19. 13
- 21. (Currently amended) Use of 1 (1R, 2S, 4R) - (-) - 2 - (2 - dimethylaminoethoxy) - 2 - phenyl - 1, 7, 7 - trimethyl - b2 icyclo[2.2.1]heptane-fumarate in the treatment of neuroprotective 3 disorders, for the reduction of the consequences of acute ischemic or traumatic brain and spinal damages, especially the various types 5 of stroke or cerebral vasospasm, severe brain vessel occlusion, neuronal loss and its functional consequences in the case of head and spinal injuries caused by accidents, for the treatment of 8 chronic neurodegenerative diseases or for the treatment of motoneuron disease (ALS), sclerosis multiplex or CreutzfeldJakob 10 disease indications according to Claims 16-19. 11

22. (Currently amended) Neuroprotective method of
treatment which comprises administering to the patient in need of
such treatment a compound of the general Formula I or a
pharmaceutically acceptable acid addition salt thereof as defined
in claim 1, perferably preferably (1R,2S,4R)-(-)-2(2-dimethylaminoethoxy)-2-phenyl-1,7,7-trimethylbicyclo[2.2.1]hepta
ne of the Formula II or a pharmaceutically acceptable acid addition
salt thereof in a therapeuticly therapeutically active amount.